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§ 436.106 Microbiological turbidimetric assay.

Using the sample solution prepared as described in the section for the particular antibiotic to be tested, proceed as described in paragraphs (a), (b), and (c) of this section.

(a) Preparation of working standard stock solutions and standard response line solutions. For each antibiotic listed in the table in this paragraph, select the working standard, drying conditions, solvent(s), concentrations, and storage time for the standard solutions and proceed as follows: If necessary, dry

the working standard as described in §436.200; dissolve and dilute an accurately weighed portion to the proper concentration for the working standard stock solution. Store the working standard stock solution under refrigeration and do not use longer than the recommended storage time. Prepare the proper concentrations for the standard response line solutions by further diluting an aliquot of the working standard stock solution. The reference concentration of the assay is the mid concentration of the standard response line.

		Workin	Working standard stock solutions	SI		Standard re	Standard response line concentrations
Antibiotic	Drying conditions (method number as listed in §436.200)	Initial solvent	Diluent (solution num- ber as listed in §436.101(a))	Final concentra- tion units or milli- grams per milli- liter	Storage time under refrigera- tion	Diluent (solution number as listed in § 436.101(a))	Final concentrations—units or micrograms of antibiotic activity per milliliter
Amikacin	Not dried6		Distilled water	1 mg	2 weeks Use same day	Distilled water	8.0, 8.9, 10.0, 11.2, 12.5 µg. 0.030, 0.043, 0.060, 0.085, 0.120
Capreomycin	5	Distilled water		1 mg	7 days	Distilled water	se line simultaneously wample solution.)
Chloramphenicol	Not dried	Ethyl alcohol (10,000	Distilled water	1 mg	1 month	Distilled water	ид. 2.00, 2.24, 2.50, 2.80, 3.12 µg.
Chlortetracycline	Not dried	rig. bd.	0.01N HCI	1 mg	4 days	Distilled water	0.048, 0.054, 0.060, 0.067, 0.075
Cycloserine			Distilled water	1 mg	1 month	Distilled water	40.0, 44.5, 50.0, 56.0, 62.5 μg. 0.080, 0.089, 0.100, 0.112, 0.125
Dihydrostreptomycin Doxycycline	5 Not dried		Distilled water	1 mg	30 days5 days	Distilled water	μg. 24.0, 26.8, 30.0, 33.5, 37.5 μg. 0.080, 0.089, 0.100, 0.112, 0.125
Gramicidin	1		alcohol U.S.P. XX	1 mg	30 days	alcohol U.S.P.	нд. 0.032, 0.0356, 0.040, 0.0448, 0.050 нд
KanamycinLincomycin	Not driedNot dried		Distilled water	1 mg	1 month	Distilled water	8.0, 8.9, 10.0, 11.2, 12.5 μg. 0.400, 0.447, 0.500, 0.559, 0.625
Meclocycline	Not dried		13	1 mg	Use same day	Distilled water	μg. 0.048, 0.054, 0.06, 0.067, 0.075
Methacycline	1		Distilled water	1 mg	7 days	Distilled water	μg. 0.048, 0.054, 0.060, 0.067, 0.075
Oxytetracycline	Not dried		0.1 N HCI	1 mg	4 days	Distilled water	ну. 0.192, 0.215, 0.240, 0.268, 0.300
Rolitetracycline	1		Distilled water	1 mg	1 day	Distilled water	hg. 0.192, 0.215, 0.240, 0.268, 0.300
Spectinomycin Streptomycin Tetracycline	Not dried1 Not dried		Distilled water Distilled water 0.1N HCI	1 mg	1 month	Distilled water Distilled water Distilled water	24.0, 26.8, 30.0, 33.5, 37.5 μg. 24.0, 26.8, 30.0, 33.5, 37.5 μg. 0.192, 0.215, 0.240, 0.268, 0.300
Tobramycin	Not dried1		Distilled water	1 mg	2 weeks	Distilled water Distilled water	μy. 2.00, 2.236, 2.5, 2.795, 3.125 μg. 20.0, 22.25, 25.0, 28.0, 31.25 μg.

¹ Use sterile equipment for all stages of this assay.
² The gramicidin working standard and the gramicidin standard response line concentrations are used for the assay of tyrothricin.

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(b) Procedure for assay. For each antibiotic listed in the table in this paragraph, select the test organism (as listed in §436.103(a)), nutrient broth (as listed by medium number in §436.102(b)), and suggested inoculum and proceed as follows: Place 1.0 milliliter (or 0.1 milliliter in the case of gramicidin and tyrothricin) of each concentration of the standard response line (prepare as described in paragraph (a) of this section) and of the sample solution in each set of three replicate tubes (as described in \$436.100(b)(1)). Fifteen tubes are used for the fivepoint standard response line and three for each sample. To each tube add 9 milliliters of the inoculated broth and place immediately in a water bath at the appropriate temperature for 2 to 4

hours. The exact length of the incubation period should be determined by observation of growth in the reference concentration tube of the standard. Remove the tubes from the water bath and add 0.5 milliliter of a 12-percent formaldehyde solution to each tube. Determine the absorbance value of each tube in a suitable photoelectric colorimeter, at a wavelength of 530 millimicrons. Set the instrument at zero absorbance with an uninoculated blank composed of the same amounts of nutrient broth and formaldehyde used in the assay.

NOTE: The amount of working standard and sample solutions may be reduced as long as all other solutions used are reduced proportionately.

Antibiotic	Test orga- nism	Medium (nutrient broth)	Suggested volume of standard-ized inoculum to be added to each 100 milliliters of medium (nutrient broth)	Incubation tempera- ture
Australia			0.4	00.07.5
Amikacin Candicidin ¹	l A	3 13	0.1 0.2	36–37.5 27–29
Capreomycin		3	0.05	36–37.5
Chloramphenicol	l j	3	0.03	36–37.5
Chlortetracycline	l ă	3	0.1	36–37.5
Cycloserine	A	3	0.4	36–37.5
Demeclocycline	A	3	0.1	36–37.5
Dihydrostreptomycin	l 1	3	0.1	36-37.5
Doxycycline	l A	3	0.1	36-37.5
Gramicidin	l ĸ	3	1.0	36-37.5
Kanamycin	A	3	0.2	36-37.5
Lincomycin	A	3	0.1	36-37.5
Meclocycline	A	3	0.2	36-37.5
Methacycline	A	3	0.1	36-37.5
Oxytetracycline	A	3	0.1	36-37.5
Rolitetracycline	A	3	0.1	36–37.5
Spectinomycin	J	3	0.1	36-37.5
Streptomycin	1	3	0.1	36-37.5
Tetracycline	A	3	0.1	36–37.5
Tobramycin	A	3	0.15	36–37.5
Troleandomycin	1	3	0.1	36–37.5
Tyrothricin	K	3	1.0	36–37.5

¹Use sterile equipment for all stages of this assay.

$$L = \frac{3a + 2b + c - e}{5}$$

$$H = \frac{3e + 2d + c - a}{5}$$

L=Calculated absorbance value for the lowest concentration of the standard response line.

H=Calculated absorbance value for the highest concentration of the standard response line.

where: *L*=Calculated absorbance value for the lowest concentration of the standard response line.

- a. b. c. d. e=Average absorbance values for each concentration of the standard response line, lowest to the highest, respectively.
- (c) Estimation of potency. To prepare the standard response line, plot the average absorbance values for each concentration of the standard response on one-cycle semilogarithmic graph paper with the absorbance values on the arithmetric scale and concentrations on the logarithmic scale. The response line is drawn either through these points by inspection or through points plotted for highest and lowest absorbance values obtained by means of the following equations.

To estimate the potency of the sample, average the absorbance values for the sample and determine the antibiotic concentration from the standard response line. Multiply the concentration by the appropriate dilution factor to obtain the antibiotic content of the sample.

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Subpart E—General Chemical **Tests for Antibiotics**

§436.200 Loss on drying.

Use the method specified in the individual section for each antibiotic.

(a) Method 1. In an atmosphere of about 10 percent relative humidity, grind the sample, if necessary, to obtain a fine powder. When tablets, troches, or capsules are to be tested, use four tablets, troches, or capsules in preparing the sample. Transfer about 100 milligrams of the sample to a tared weighing bottle equipped with a ground-glass stopper. Weigh the bottle and place it in a vacuum oven, tilting the stopper on its side so that there is no closure during the drying period. Dry at a temperature of 60° C. and a pressure of 5 millimeters of mercury or less for 3 hours. At the end of the drying period, fill the vacuum oven with air dried by passing it through a drying agent such as sulfuric acid or silica gel.

Replace the stopper and place the weighing bottle in a desiccator over a desiccating agent, such as phosphorous pentoxide or silica gel, allow to cool to room temperature, and reweigh. Calculate the percent of loss.

(b) Method 2. Proceed as directed in paragraph (a) of this section, except use a tared weighing bottle or weighing tube equipped with a capillary-tube stopper, the capillary having an inside diameter of 0.20-0.25 millimeter, and place it in a vacuum oven without removing the stopper.

(c) Method 3. Proceed as directed in paragraph (a) of this section, except dry the sample at a temperature of 110° C. and a pressure of 5 millimeters of

mercury or less for 3 hours.

(d) Method 4. Proceed as directed in paragraph (a) of this section, except dry the sample at a temperature of 40° C. and a pressure of 5 millimeters of mercury or less for 2 hours.

(e) Method 5. Proceed as directed in paragraph (a) of this section, except dry the sample at a temperature of 100° C. and a pressure of 5 millimeters of mercury or less for 4 hours.

(f) Method 6. Proceed as directed in paragraph (a) of this section, except dry the sample at a temperature of 40° C. and a pressure of 5 millimeters of mercury or less for 3 hours.

(g) Method 7. Proceed as directed in paragraph (a) of this section, except dry the sample at a temperature of 25° C. and a pressure of 5 millimeters of

mercury or less for 4 hours.

- (h) Method 8. Proceed as directed in paragraph (a) of this section, except transfer approximately 300 milligrams of the sample to a tared weighing bottle equipped with a ground-glass stopper; dry the sample at a temperature of 25 ° C and a pressure of 5 millimeters of mercury or less for 4 hours, and then dry the sample at a temperature of 100 °C and a pressure of 5 millimeters of mercury or less for 3 additional hours.
- *Method 9.* Use a suitable thermogravimetric apparatus prepared for vacuum operation. Rapidly and thoroughly grind a portion of the sample and promptly transfer 5 to 10 milligrams to the sample pan. Place the system under vacuum and allow it to come to equilibrium before heating. Obtain an accurate sample weight and